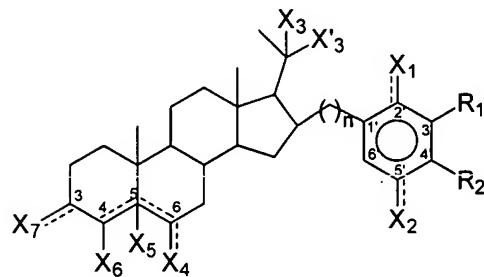


AMENDMENTS TO THE CLAIMS

1. (Currently amended) A compound having the structural formula IB or a pharmaceutically acceptable salt thereof,



formula IB

wherein X₁, X₂, R₁ and R₂ are independently selected from the group comprising consisting of oxo, hydrogen, hydroxyl, oxyalkyl, alkyl, alkenyl, alkynyl, alkyloxy, alkyloxyalkyl, alkylthioalkyl, alkoxycarbonyl, alkylthiocarbonyl, alkanoyl, cycloalkylalkyl, cycloalkylcarbonyl, cycloalkylalkanoyl, cycloalkylthiocarbonyl, cycloalkylalkoxycarbonyl, cycloalkylalkoxythiocarbonyl, cycloalkylcarbonyloxyalkyl, cycloalkylcarbonyloxyalkyl, silyloxyalkyl, aralkyl, arylalkenyl, arylcarbonyl, aryloxycarbonyl, arylthiocarbonyl, aralkoxycarbonyl, arylalkylthiocarbonyl, aryloxyalkyl, arylthioalkyl, haloalkyl, hydroxyalkyl, aralkanoyl, aroyl, aryloxycarbonylalkyl, aryloxyalkanoyl, carboxyl, alkenylcarbonyl, alkynylcarbonyl, Het¹, Het¹alkyl, Het¹oxyalkyl, Het¹aryl, Het¹aralkyl, Het¹cycloalkyl, Het¹alkoxycarbonyl, Het¹alkylthiocarbonyl, Het¹oxycarbonyl, Het¹thiocarbonyl, Het¹alkanoyl, Het¹aralkanoyl, Het¹aryloxyalkyl, Het¹alkyloxyalkyl, Het¹arylthioalkyl, Het¹aryloxycarbonyl, Het¹aralkoxycarbonyl, Het¹aroyl, Het¹oxyalkylcarbonyl, Het¹alkyloxyalkylcarbonyl, Het¹aryloxyalkylcarbonyl, Het¹carbonyloxyalkyl, Het¹alkylcarbonyloxyalkyl, Het¹aralkylcarbonyloxyalkyl, Het²alkyl, Het²oxyalkyl, Het²alkyloxyalkyl, Het²aralkyl, Het²carbonyl, Het²oxycarbonyl, Het²thiocarbonyl, Het²alkanoyl, Het²alkylthiocarbonyl, Het²alkoxycarbonyl, Het²aralkanoyl, Het²aralkoxycarbonyl, Het²aryloxycarbonyl, Het²aroyl, Het²aryloxyalkyl, Het²arylthioalkyl, Het²oxyalkylcarbonyl, Het²alkyloxyalkylcarbonyl, Het²aryloxyalkylcarbonyl, Het²carbonyloxyalkyl, Het²alkylcarbonyloxyalkyl, Het²aralkylcarbonyloxyalkyl, cyano, CR³=NR⁴, CR³=N(OR⁴), aminocarbonyl, aminoalkanoyl, aminoalkyl, optionally-unsubstituted or substituted by one or more substituents independently selected from the group comprising-consisting of alkyl, aralkyl, aryl, Het¹, Het², cycloalkyl, alkyloxy, alkyloxycarbonyl, carboxyl, aminocarbonyl,

mono- or di(alkyl)aminocarbonyl, aminosulfonyl, alkylS(=O)_t, hydroxy, cyano, halogen or amino, optionally unsubstituted, mono- or disubstituted, wherein the substituents are independently selected from the group comprising consisting of alkyl, aryl, aralkyl, aryloxy, arylamino, arylthio, aryloxyalkyl, arylaminoalkyl, aralkoxy, alkylthio, alkoxy, aryloxyalkoxy, arylaminoalkoxy, aralkylamino, aryloxyalkylamino, arylaminoalkylamino, arylthioalkoxy, arylthioalkylamino, aralkylthio, aryloxyalkylthio, arylaminoalkylthio, arylthioalkylthio, alkylamino, cycloalkyl, cycloalkylalkyl, Het¹, Het², Het¹alkyl, Het²alkyl, Het¹amino, Het²amino, Het¹alkylamino, Het²alkylamino, Het¹thio, Het²thio, Het¹alkylthio, Het²alkylthio, Het¹oxy and Het²oxy, OR³, SR³, SO₂NR³R⁴, SO₂N(OH)R³, CN, CR³=NR⁴, S(O)R³, SO₂R³, CR³=N(OR⁴), N₃, NO₂, NR³R⁴, N(OH)R³, C(O)R³, C(S)R³, CO₂R³, C(O)SR³, C(O)NR³R⁴, C(S)NR³R⁴, C(O)N(OH)R⁴, C(S)N(OH)R³, NR³C(O)R⁴, NR³C(S)R⁴, N(OH)C(O)R⁴, N(OH)C(S)R³, NR³CO₂R⁴, NR³C(O)NR⁴R⁵, and NR³C(S)NR⁴R⁵, N(OH)CO₂R³, NR³C(O)SR⁴, N(OH)C(O)NR³R⁴, N(OH)C(S)NR³R⁴, NR³C(O)N(OH)R⁴, NR³C(S)N(OH)R⁴, NR³SO₂R⁴, NHSO₂NR³R⁴, NR³SO₂NHR⁴, P(O)(OR³)(OR⁴), wherein t is an integer between 1 and 2 and R³, R⁴ and R⁵ are each independently selected from the group comprising consisting of hydrogen, hydroxyl, alkyl, alkenyl, alkynyl, aminoalkyl, aminoaryl, alkylcarbonylamino, arylcarbonylamino, alkylthiocarbonylamino and arylthiocarbonylamino;

wherein X₃ participates together with X₃' to in an oxo functional group, or wherein X₃ is selected from the group comprising consisting of hydrogen, hydroxyl, sulfur, oxyalkyl, oxycarbonyl, alkyl, Het¹alkyl, alkenyl, alkynyl, aminoalkyl, aminoacyl, alkylcarbonylamino, alkylthiocarbonylamino, Het¹, glycosyl, thio derivatives thereof, amino derivatives thereof, hydroxyl-protected derivatives thereof, alkyloxycarbonyl, optionally unsubstituted or substituted by one or more substituents independently selected from the group comprising consisting of alkyl, aralkyl, aryl, Het¹, Het², cycloalkyl, alkyloxy, alkyloxycarbonyl, carboxyl and aminocarbonyl; and X₃ is selected from the group comprising consisting of hydrogen, alkyl, aryl, Het¹, glycosyl, thio derivatives thereof, amino derivatives thereof, hydroxyl-protected derivatives thereof, aralkyl, and optionally unsubstituted or substituted by one or more substituents independently selected from the group comprising consisting of alkyl, aralkyl, aryl, Het¹, Het², cycloalkyl, alkyloxy, alkyloxycarbonyl, carboxyl, aminocarbonyl, mono- or di(alkyl)aminocarbonyl, aminosulfonyl, alkylS(=O)_t, hydroxy, cyano, halogen or amino, optionally unsubstituted, mono- or disubstituted, wherein the substituents are independently selected from the group comprising consisting of alkyl, aryl, aralkyl, aryloxy, arylamino, arylthio, aryloxyalkyl, arylaminoalkyl, aralkoxy, alkylthio, alkoxy, aryloxyalkoxy, arylaminoalkoxy, aralkylamino,

aryloxyalkylamino, arylaminoalkylamino, arylthioalkoxy, arylthioalkylamino, aralkylthio, aryloxyalkylthio, arylaminoalkylthio, arylthioalkylthio, alkylamino, cycloalkyl and cycloalkylalkyl;

wherein X_4 and X_7 are independently selected from the group comprising-consisting of hydrogen, halogen, oxygen, oxo, carbonyl, thiocarbonyl, hydroxyl, alkyl, aryl, Het^1 , glycosyl, thio derivatives thereof, amino derivatives thereof, hydroxyl-protected derivatives thereof, Het^1 alkyl, Het^1 aryl, alkenyl, alkynyl, hydroxyalkyl, hydroxycarbonyl, hydroxycarbonylalkyl, hydroxycarbonylaryl, hydroxycarbonyloxyalkyl and hydroxycarbonyloxyaryl; aminocarbonyl, mono- or di(alkyl)aminocarbonyl, aminosulfonyl, alkylS(=O)_t, hydroxy, aminoalkyl, aminoaryl, cyano, halogen or amino, unsubstituted, optionally mono- or disubstituted, wherein the substituents are independently selected from the group comprising-consisting of alkyl, aryl, aralkyl, aryloxy, arylamino, arylthio, aryloxyalkyl, arylaminoalkyl, aralkoxy, alkoxy, aryloxyalkoxy, aylaminoalkoxy, aralkylamino, aryloxyalkylamino, arylaminoalkylamino, arylthioalkoxy, arylthioalkylamino, aralkylthio, aryloxyalkylthio, arylaminoalkylthio, arylthioalkylthio, alkylamino, Het^1 , Het^2 , alkyloxycarbonyl, carboxyl, aminocarbonyl, cycloalkyl and cycloalkylalkyl;

wherein X_5 participates in a double bond between the carbon atoms in position 4 and 5 or between carbon atoms in position 5 and 6, and X_6 is independently selected from the group comprising-consisting of hydrogen, hydroxyl and hydroxyalkyl, or wherein X_5 and X_6 are independently selected from the group comprising-consisting of halogen, hydrogen, hydroxyl, hydroxyalkyl, aminoalkyl, aminoaryl, optionally-unsubstituted or substituted by one or more substituents independently selected from the group comprising-consisting of alkyl, aralkyl, aryl, Het^1 , Het^2 , cycloalkyl, alkyloxy, alkyloxycarbonyl, carboxyl, aminocarbonyl, and

wherein n is an integer between 0 and 10,

provided that when X_6 and X_4 are H, when X_5 participates in a double bond between the carbon atoms in position 5 and 6, when X_3 participates together with X_3' in an oxo functional group, when n is zero and X_1 , X_2 , R_1 and R_2 are H, X_7 is not hydroxyl.

2. (Currently amended) A-The compound according to claim 1,

wherein X_1 , X_2 , R_1 and R_2 are independently selected from the group comprising-consisting of oxo, hydrogen, hydroxyl, oxyalkyl, alkyl, alkenyl, alkynyl, alkyloxy, alkyloxalkyl, alkylthioalkyl, alkoxy carbonyl, alkylthiocarbonyl, alkanoyl, cycloalkylalkyl, cycloalkylcarbonyl,

cycloalkylalkanoyl, cycloalkylthiocarbonyl, cycloalkylalkoxycarbonyl,
cycloalkylalkoxythiocarbonyl, cycloalkylthioalkyl, alkylcarbonyloxyalkyl,
cycloalkylcarbonyloxyalkyl, silyloxyalkyl, aralkyl, arylalkenyl, arylcarbonyl, aryloxycarbonyl,
arylthiocarbonyl, aralkoxycarbonyl, arylalkylthiocarbonyl, aryloxyalkyl, arylthioalkyl, haloalkyl,
hydroxyalkyl, aralkanoyl, aroyl, aryloxycarbonylalkyl, aryloxyalkanoyl, carboxyl, alkenylcarbonyl,
alkynylcarbonyl, Het¹, Het¹alkyl, Het¹oxyalkyl, Het¹aryl, Het¹aralkyl, Het¹cycloalkyl,
Het¹alkoxycarbonyl, Het¹alkylthiocarbonyl, Het¹oxycarbonyl, Het¹thiocarbonyl, Het¹alkanoyl,
Het¹aralkanoyl, Het¹aryloxyalkyl, Het¹alkyloxyalkyl, Het¹arylthioalkyl, Het¹aryloxycarbonyl,
Het¹aralkoxycarbonyl, Het¹aroyl, Het¹oxyalkylcarbonyl, Het¹alkyloxyalkylcarbonyl,
Het¹aryloxyalkylcarbonyl, Het¹carbonyloxyalkyl, Het¹alkylcarbonyloxyalkyl,
Het¹aralkylcarbonyloxyalkyl, Het²alkyl, Het²oxyalkyl, Het²alkyloxyalkyl, Het²aralkyl, Het²carbonyl,
Het²oxycarbonyl, Het²thiocarbonyl, Het²alkanoyl, Het²alkylthiocarbonyl, Het²alkoxycarbonyl,
Het²aralkanoyl, Het²aralkoxycarbonyl, Het²aryloxycarbonyl, Het²aroyl, Het²aryloxyalkyl,
Het²arylthioalkyl, Het²oxyalkylcarbonyl, Het²alkyloxyalkylcarbonyl, Het²aryloxyalkylcarbonyl,
Het²carbonyloxyalkyl, Het²alkylcarbonyloxyalkyl, Het²aralkylcarbonyloxyalkyl, cyano, CR³=NR⁴,
CR³=N(OR⁴), aminocarbonyl, aminoalkanoyl, aminoalkyl, optionally-unsubstituted or substituted
by one or more substituents independently selected from the group comprising-consisting of
alkyl, aralkyl, aryl, Het¹, Het², cycloalkyl, alkyloxy, alkyloxycarbonyl, carboxyl, aminocarbonyl,
mono- or di(alkyl)aminocarbonyl, aminosulfonyl, alkylS(=O)_t, hydroxy, cyano, halogen or amino,
optionally-unsubstituted, mono- or disubstituted, wherein the substituents are independently
selected from the group comprising-consisting of alkyl, aryl, aralkyl, aryloxy, arylamino, arylthio,
aryloxyalkyl, arylaminoalkyl, aralkoxy, alkylthio, alkoxy, aryloxyalkoxy, aylaminoalkoxy,
aralkylamino, aryloxyalkylamino, arylaminoalkylamino, arylthioalkoxy, arylthioalkylamino,
aralkylthio, aryloxyalkylthio, arylaminoalkylthio, arylthioalkylthio, alkylamino, cycloalkyl,
cycloalkylalkyl, Het¹, Het², Het¹alkyl, Het²alkyl, Het¹amino, Het²amino, Het¹alkylamino,
Het²alkylamino, Het¹thio, Het²thio, Het¹alkylthio, Het²alkylthio, Het¹oxy-and, Het²oxy, OR³, SR³,
SO₂NR³R⁴, SO₂N(OH)R³, CN, CR³=NR⁴, S(O)R³, SO₂R³, CR³=N(OR⁴), N₃, NO₂, NR³R⁴,
N(OH)R³, C(O)R³, C(S)R³, CO₂R³, C(O)SR³, C(O)NR³R⁴, C(S)NR³R⁴, C(O)N(OH)R⁴,
C(S)N(OH)R³, NR³C(O)R⁴, NR³C(S)R⁴, N(OH)C(O)R⁴, N(OH)C(S)R³, NR³CO₂R⁴,
NR³C(O)NR⁴R⁵, and—NR³C(S)NR⁴R⁵, N(OH)CO₂R³, NR³C(O)SR⁴, N(OH)C(O)NR³R⁴,
N(OH)C(S)NR³R⁴, NR³C(O)N(OH)R⁴, NR³C(S)N(OH)R⁴, NR³SO₂R⁴, NHSO₂NR³R⁴,
NR³SO₂NHR⁴, P(O)(OR³)(OR⁴), wherein t is an integer between 1 and 2 and R³, R⁴ and R⁵ are
each independently selected from the group comprising-consisting of hydrogen, hydroxyl, alkyl,

alkenyl, alkynyl, aminoalkyl, aminoaryl, alkylcarbonylamino, arylcarbonylamino alkylthiocarbonylamino and arylthiocarbonylamino;

wherein X_3 participates together with X_3' to in an oxo functional group, or wherein X_3 is selected from the group comprising consisting of hydrogen, hydroxyl, sulfur, oxyalkyl, oxycarbonyl, alkyl, Het¹alkyl, alkenyl, alkynyl, aminoalkyl, aminoacyl, alkylcarbonylamino, alkylthiocarbonylamino, Het¹, glucosyl, fructosyl, galactosyl, mannosyl, ribosyl, ribulosyl, xylulosyl, erythrosyl, erythrulosyl, rhamnosyl, threosyl, sorbosyl, psicosyl, tagatosyl, fucosyl, arabinosyl, xylofuranosyl, lyxosyl, talosyl, psicosyl, idosyl, gulosyl, altrosyl, allosyl, mannoheptulosyl, sedoheptulosyl, abequosyl, isomaltosyl, kojibiosyl, laminarabiosyl, nigerosyl, primeverosyl, rutinosyl, tyvelosyl, maltosyl, lactosyl, sucrosyl, cellobiosyl, trehalosyl, gentiobiosyl, melibiosyl, turanosyl, sophorosyl, isosucrosyl, raffinosyl, gentianosyl, 2-amino-2-deoxy glucosyl, 2-acetamido-2-deoxy-glucosyl, 2-amino-2-deoxy galactosyl, 2-acetamido-2-deoxy-galactosyl, 2-amino-2-deoxy mannosyl, 2-acetamido-2-deoxy-mannosyl, 2-amino-1,3-cyclohexanediol, L or D isomers thereof, α or β form thereof, pyranose or furanose form thereof, combination thereof, deoxy derivatives thereof, hydroxyl-protected acetate derivatives thereof, amino derivatives thereof, thio derivatives thereof, di-, tri-, oligo- and polysaccharide thereof, alkyloxycarbonyl, optionally unsubstituted or substituted by one or more substituents independently selected from the group comprising consisting of alkyl, aralkyl, aryl, Het¹, Het², cycloalkyl, alkyloxy, alkyloxycarbonyl, carboxyl and aminocarbonyl; and X_3 is selected from the group comprising consisting of hydrogen, alkyl, aryl, Het¹, glucosyl, fructosyl, galactosyl, mannosyl, ribosyl, ribulosyl, xylulosyl, erythrosyl, erythrulosyl, rhamnosyl, threosyl, sorbosyl, psicosyl, tagatosyl, fucosyl, arabinosyl, xylofuranosyl, lyxosyl, talosyl, psicosyl, idosyl, gulosyl, altrosyl, allosyl, mannoheptulosyl, sedoheptulosyl, abequosyl, isomaltosyl, kojibiosyl, laminarabiosyl, nigerosyl, primeverosyl, rutinosyl, tyvelosyl, maltosyl, lactosyl, sucrosyl, cellobiosyl, trehalosyl, gentiobiosyl, melibiosyl, turanosyl, sophorosyl, isosucrosyl, raffinosyl, gentianosyl, 2-amino-2-deoxy glucosyl, 2-acetamido-2-deoxy-glucosyl, 2-amino-2-deoxy galactosyl, 2-acetamido-2-deoxy-galactosyl, 2-amino-2-deoxy mannosyl, 2-acetamido-2-deoxy-mannosyl, 2-amino-1,3-cyclohexanediol, L or D isomers thereof, α or β form thereof, pyranose or furanose form thereof, combination thereof, deoxy derivatives thereof, hydroxyl-protected acetate derivatives thereof, amino derivatives thereof, thio derivatives thereof, di-, tri-, oligo- and polysaccharide thereof, aralkyl, and optionally unsubstituted or substituted by one or more substituents independently selected from the group comprising consisting of alkyl, aralkyl, aryl, Het¹, Het², cycloalkyl, alkyloxy, alkyloxycarbonyl, carboxyl, aminocarbonyl, mono- or

di(alkyl)aminocarbonyl, aminosulfonyl, alkylS(=O)_t, hydroxy, cyano, halogen or amino, optionally unsubstituted, mono- or disubstituted, wherein the substituents are independently selected from the group comprising consisting of alkyl, aryl, aralkyl, aryloxy, arylamino, arylthio, aryloxyalkyl, arylaminoalkyl, aralkoxy, alkylthio, alkoxy, aryloxyalkoxy, aylaminoalkoxy, aralkylamino, aryloxyalkylamino, arylaminoalkylamino, arylthioalkoxy, arylthioalkylamino, aralkylthio, aryloxyalkylthio, arylaminoalkylthio, arylthioalkylthio, alkylamino, cycloalkyl and cycloalkylalkyl;

wherein X₄ and X₇ are independently selected from the group comprising consisting of hydrogen, oxygen, halogen, oxo, carbonyl, thiocarbonyl, hydroxyl, alkyl, aryl, Het¹, glucosyl, fructosyl, galactosyl, mannosyl, ribosyl, ribulosyl, xylulosyl, erythrosyl, erythrulosyl, rhamnosyl, threosyl, sorbosyl, psicosyl, tagatosyl, fucosyl, arabinosyl, xylofuranosyl, lyxosyl, talosyl, psicosyl, idosyl, gulosyl, altrosyl, allosyl, mannoheptulosyl, sedoheptulosyl, abequosyl, isomaltosyl, kojibiosyl, laminarabiosyl, nigerosyl, primeverosyl, rutinosyl, tyvelosyl, maltosyl, lactosyl, sucrosyl, cellobiosyl, trehalosyl, gentiobiosyl, melibiosyl, turanosyl, sophorosyl, isosucrosyl, raffinosyl, gentianosyl, 2-amino-2-deoxy glucosyl, 2-acetamido-2-deoxy-glucosyl, 2-amino-2-deoxy galactosyl, 2-acetamido-2-deoxy-galactosyl, 2-amino-2-deoxy mannosyl, 2-acetamido-2-deoxy-mannosyl, 2-amino-1,3-cyclohexanediol, L or D isomers thereof, α or β form thereof, pyranose or furanose form thereof, combination thereof, deoxy derivatives thereof, hydroxyl-protected acetate derivatives thereof, amino derivatives thereof, thio derivatives thereof, di-, tri-, oligo- and polysaccharide thereof; Het¹alkyl, Het¹aryl, alkenyl, alkynyl, hydroxyalkyl, hydroxycarbonyl, hydroxycarbonylalkyl, hydroxycarbonylaryl, hydroxycarbonyloxyalkyl, and hydroxycarbonyloxyaryl; aminocarbonyl, mono- or di(alkyl)aminocarbonyl, aminosulfonyl, alkylS(=O)_t, hydroxy, aminoalkyl, aminoaryl, cyano, halogen or amino, optionally unsubstituted, mono- or disubstituted wherein the substituents are independently selected from the group comprising consisting of alkyl, aryl, aralkyl, aryloxy, arylamino, arylthio, aryloxyalkyl, arylaminoalkyl, aralkoxy, alkylthio, alkoxy, aryloxyalkoxy, aylaminoalkoxy, aralkylamino, aryloxyalkylamino, arylaminoalkylamino, arylthioalkoxy, arylthioalkylamino, aralkylthio, aryloxyalkylthio, arylaminoalkylthio, arylthioalkylthio, alkylamino, Het¹, Het², alkyloxycarbonyl, carboxyl, aminocarbonyl, cycloalkyl and cycloalkylalkyl;

wherein X₅ participates to in a double bond between the carbon atoms in position 4 and 5 or between carbon atoms in positions 5 and 6, and X₆ is independently selected from the group comprising consisting of hydrogen, hydroxyl and hydroxyalkyl, or

wherein X_5 and X_6 are independently selected from the group comprising consisting of halogen, hydrogen, hydroxyl, hydroxyalkyl, aminoalkyl, aminoaryl, optionally unsubstituted or substituted by one or more substituents independently selected from the group comprising consisting of alkyl, aralkyl, aryl, Het^1 , Het^2 , cycloalkyl, alkyloxy, alkyloxycarbonyl, carboxyl, aminocarbonyl, and

wherein n is an integer between 0 and 10.

3. (Currently amended) A The compound according to claim 1-~~or~~ 2,

wherein X_1 , X_2 , R_1 and R_2 ~~is~~ are selected from the group comprising consisting of hydrogen, hydroxyl, oxyalkyl, oxo, alkyl, alkenyl, alkynyl, alkyloxy, alkyloxyalkyl, alkylthioalkyl, alkoxy carbonyl, alkylthiocarbonyl, alkanoyl, cycloalkylalkyl, cycloalkylcarbonyl, cycloalkylalkanoyl, cycloalkylthiocarbonyl, cycloalkylalkoxy carbonyl, cycloalkylalkoxythiocarbonyl, cycloalkylthioalkyl, alkylcarbonyloxyalkyl, cycloalkylcarbonyloxyalkyl, silyloxyalkyl, aralkyl, arylalkenyl, arylcarbonyl, aryloxycarbonyl, arylthiocarbonyl, aralkoxycarbonyl, arylalkylthiocarbonyl, aryloxyalkyl, arylthioalkyl, haloalkyl, hydroxyalkyl, aralkanoyl, aroyl, aryloxycarbonylalkyl, aryloxyalkanoyl, carboxyl, alkenylcarbonyl and alkynylcarbonyl;

wherein X_3 participates together with X_3' ~~to~~ in an oxo functional group, or wherein X_3 is selected from the group comprising consisting of hydrogen, hydroxyl, sulfur, oxyalkyl, oxycarbonyl alkyl, Het^1 alkyl, alkenyl, alkynyl, aminoalkyl, aminoacyl, alkylcarbonylamino, alkylthiocarbonylamino, Het^1 , glucosyl, fructosyl, galactosyl, mannosyl, ribosyl, ribulosyl, xylulosyl, erythrosyl, erythrulosyl, rhamnosyl, threosyl, sorbosyl, psicosyl, tagatosyl, fucosyl, arabinosyl, xylofuranosyl, lyxosyl, talosyl, psicosyl, idosyl, gulosyl, altrosyl, allosyl, mannoheptulosyl, sedoheptulosyl, abequosyl, isomaltosyl, kojibiosyl, laminarabiosyl, nigerosyl, primeverosyl, rutinosyl, tyvelosyl, maltosyl, lactosyl, sucrosyl, cellobiosyl, trehalosyl, gentiobiosyl, melibiosyl, turanosyl, sophorosyl, isosucrosyl, raffinosyl, gentianosyl, 2-amino-2-deoxy glucosyl, 2-acetamido-2-deoxy-glucosyl, 2-amino-2-deoxy galactosyl, 2-acetamido-2-deoxy-galactosyl, 2-amino-2-deoxy mannosyl, 2-acetamido-2-deoxy-mannosyl, 2-amino-1,3-cyclohexanediol, L or D isomers thereof, α or β form thereof, pyranose or furanose form thereof, combination thereof, deoxy derivatives thereof, hydroxyl-protected acetate derivatives thereof, amino derivatives thereof, thio derivatives thereof, disaccharide thereof, trisaccharide thereof, oligosaccharide and polysaccharide thereof, alkyloxycarbonyl optionally unsubstituted

or substituted by one or more substituents independently selected from the group comprising consisting of alkyl, aralkyl, aryl, Het¹, Het², cycloalkyl, alkyloxy, alkyloxycarbonyl, carboxyl and aminocarbonyl; and X₃ is selected from the group comprising consisting of hydrogen, alkyl, aryl, aralkyl, Het¹, glucosyl, fructosyl, galactosyl, mannosyl, ribosyl, ribulosyl, xylulosyl, erythrosyl, erythrulosyl, rhamnosyl, threosyl, sorbosyl, psicosyl, tagatosyl, fucosyl, arabinosyl, xylofuranosyl, lyxosyl, talosyl, psicosyl, idosyl, gulosyl, altrosyl, allosyl, mannoheptulosyl, sedoheptulosyl, abequosyl, isomaltosyl, kojibiosyl, laminarabiosyl, nigerosyl, primeverosyl, rutinosyl, tyvelosyl, maltosyl, lactosyl, sucrosyl, cellobiosyl, trehalosyl, gentiobiosyl, melibiosyl, turanosyl, sophorosyl, isosucrosyl, raffinosyl, gentianosyl, 2-amino-2-deoxy glucosyl, 2-acetamido-2-deoxy-glucosyl, 2-amino-2-deoxy galactosyl, 2-acetamido-2-deoxy-galactosyl, 2-amino-2-deoxy mannosyl, 2-acetamido-2-deoxy-mannosyl, 2-amino-1,3-cyclohexanediol, L or D isomers thereof, α or β form thereof, pyranose or furanose form thereof, combination thereof, deoxy derivatives thereof, hydroxyl-protected acetate derivatives thereof, amino derivatives thereof, thio derivatives thereof, disaccharide thereof, trisaccharide thereof, oligosaccharide and polysaccharide thereof;

wherein X₄ and X₇ are independently selected from the group comprising consisting of hydrogen, oxygen, oxo, carbonyl, thiocarbonyl, hydroxyl, alkyl, aryl, Het¹, Het¹alkyl, Het¹aryl, alkenyl, alkynyl, hydroxyalkyl, hydroxycarbonyl, hydroxycarbonylalkyl, hydroxycarbonylaryl, hydroxycarbonyloxyalkyl, glucosyl, fructosyl, galactosyl, mannosyl, ribosyl, ribulosyl, xylulosyl, erythrosyl, erythrulosyl, rhamnosyl, threosyl, sorbosyl, psicosyl, tagatosyl, fucosyl, arabinosyl, xylofuranosyl, lyxosyl, talosyl, psicosyl, idosyl, gulosyl, altrosyl, allosyl, mannoheptulosyl, sedoheptulosyl, abequosyl, isomaltosyl, kojibiosyl, laminarabiosyl, nigerosyl, primeverosyl, rutinosyl, tyvelosyl, maltosyl, lactosyl, sucrosyl, cellobiosyl, trehalosyl, gentiobiosyl, melibiosyl, turanosyl, sophorosyl, isosucrosyl, raffinosyl, gentianosyl, 2-amino-2-deoxy glucosyl, 2-acetamido-2-deoxy-glucosyl, 2-amino-2-deoxy galactosyl, 2-acetamido-2-deoxy-galactosyl, 2-amino-2-deoxy mannosyl, 2-acetamido-2-deoxy-mannosyl, 2-amino-1,3-cyclohexanediol, L or D isomers thereof, α or β form thereof, pyranose or furanose form thereof, combinations thereof, deoxy derivatives thereof, hydroxyl-protected acetate derivatives thereof, amino derivatives thereof, thio derivatives thereof, disaccharide thereof, trisaccharide thereof, oligosaccharide and polysaccharide thereof;

wherein X₅ participates in a double bond between the carbon atoms in position 4 and 5 or between carbon atoms in positions 5 and 6, and X₆ is independently selected from the group

comprising consisting of hydrogen, hydroxyl, and hydroxyalkyl, or wherein X_5 and X_6 are independently selected from the group comprising consisting of hydrogen, hydroxyl, hydroxyalkyl, aminoalkyl, aminoaryl, optionally unsubstituted or substituted by one or more substituents independently selected from the group comprising consisting of alkyl, aralkyl, aryl, Het¹, Het², cycloalkyl, alkyloxy, alkyloxycarbonyl, carboxyl, aminocarbonyl, and

wherein n is an integer between 0 and 5.

4. (Currently amended) ~~A-The~~ compound according to ~~any of claims 1 to 3~~ claim 1,

wherein X_1 , X_2 , R_1 and R_2 ~~is-are~~ selected from the group comprising consisting of hydrogen, hydroxyl, alkyloxy, oxo and oxyalkyl,

wherein X_3 participates together with X_3' ~~to~~in an oxo functional group, or wherein X_3 is selected from the group comprising consisting of hydrogen, hydroxyl, oxyalkyl, oxycarbonyl, glucosyl, fructosyl, galactosyl, mannosyl, ribosyl, ribulosyl, xylulosyl, erythrosyl, erythrulosyl, rhamnosyl, threosyl, sorbosyl, psicosyl, tagatosyl, fucosyl, arabinosyl, 2-amino-2-deoxy glucosyl, 2-acetamido-2-deoxy-glucosyl, 2-amino-2-deoxy galactosyl, 2-acetamido-2-deoxy-galactosyl, 2-amino-2-deoxy mannosyl, 2-acetamido-2-deoxy-mannosyl, L or D isomers thereof, α or β form thereof, pyranose or furanose form thereof, combination thereof, deoxy derivatives thereof, hydroxyl-protected acetate derivatives thereof, amino derivatives thereof, thio derivatives thereof, disaccharide thereof, trisaccharide thereof, oligosaccharide and polysaccharide thereof; and X_3' is selected from the group comprising consisting of alkyl, aryl and aralkyl, glucosyl, fructosyl, galactosyl, mannosyl, ribosyl, ribulosyl, xylulosyl, erythrosyl, erythrulosyl, rhamnosyl, threosyl, sorbosyl, psicosyl, tagatosyl, fucosyl, arabinosyl, 2-amino-2-deoxy glucosyl, 2-acetamido-2-deoxy-glucosyl, 2-amino-2-deoxy galactosyl, 2-acetamido-2-deoxy-galactosyl, 2-amino-2-deoxy mannosyl, 2-acetamido-2-deoxy-mannosyl, L or D isomers thereof, α or β form thereof, pyranose or furanose form thereof, combinations thereof, deoxy derivatives thereof, hydroxyl-protected acetate derivatives thereof, amino derivatives thereof, thio derivatives thereof, disaccharide thereof, trisaccharide thereof, oligosaccharide and polysaccharide thereof;

wherein X_4 and X_7 are independently selected from the group comprising consisting of hydrogen, oxygen, oxo, hydroxyl, glucosyl, fructosyl, galactosyl, mannosyl, ribosyl, ribulosyl, xylulosyl, erythrosyl, erythrulosyl, rhamnosyl, threosyl, sorbosyl, psicosyl, tagatosyl, fucosyl, arabinosyl, 2-amino-2-deoxy glucosyl, 2-acetamido-2-deoxy-glucosyl, 2-amino-2-deoxy

galactosyl, 2-acetamido-2-deoxy-galactosyl, 2-amino-2-deoxy mannosyl, 2-acetamido-2-deoxy-mannosyl, L or D isomers thereof, α or β form thereof, pyranose or furanose form thereof, combination thereof, deoxy derivatives thereof, hydroxyl-protected acetate derivatives thereof, amino derivatives thereof, thio derivatives thereof, disaccharide thereof, trisaccharide thereof, oligosaccharide and polysaccharide thereof;

wherein X_5 and X_6 are hydrogen or wherein X_5 participates ~~to~~in a double bond between the carbon atoms in position 4 and 5, and X_6 is hydrogen, and

wherein n is an integer between 0 and 2.

5. (Currently amended) A-The compound according to ~~any of claims 1 to 4~~claim 1,

wherein X_1 , X_2 , X_3 , X_3' , X_6 , X_7 , R_1 , R_2 and n are selected from the group indicated in ~~claims 1 to 3~~claim 1; and

wherein X_4 is equal to X_5 and is selected from the group ~~comprising~~consisting of halogen, aminoalkyl, aminoaryl, ~~optionally~~unsubstituted or substituted by one or more substituents independently selected from the group ~~comprising~~consisting of alkyl, aralkyl, aryl, Het¹, Het², cycloalkyl, alkyloxy, alkyloxycarbonyl, carboxyl and aminocarbonyl, or wherein X_5 participates ~~to~~in a double bond between the carbon atoms in position 5 and 6, and X_4 is independently selected from the group ~~comprising~~consisting of hydrogen, aminoalkyl, aminoaryl, ~~optionally~~unsubstituted or substituted by one or more substituents independently selected from the group ~~comprising~~consisting of alkyl, aralkyl, aryl, Het¹, Het², cycloalkyl, alkyloxy, alkyloxycarbonyl, carboxyl and aminocarbonyl.

6. (Currently amended) A-The compound according to ~~any of claims 1 to 4~~Claim 1, wherein X_1 and X_2 are $-OMe$, wherein R_1 and R_2 are $-H$, wherein X_4 is hydrogen, wherein X_3 participates together with X_3' ~~to~~in an oxo functional group, wherein X_5 participates ~~to~~in a double bond between the carbon atoms in position 4 and 5, wherein X_6 is hydrogen, wherein X_7 is hydroxyl, glucosyl, fructosyl, galactosyl, mannosyl, fucosyl, cellobiosyl, gentiobiosyl, 2-amino-2-deoxy glucosyl, 2-acetamido-2-deoxy-glucosyl, 2-amino-2-deoxy galactosyl, 2-acetamido-2-deoxy-galactosyl, disaccharide or trisaccharide thereof; and wherein n is 0.

7. (Currently amended) A-The compound according to ~~any of claims 1 to 4~~claim 1, wherein X_1 and X_2 are $-OMe$, wherein R_1 and R_2 are $-H$, wherein X_3 is hydrogen, hydroxyl, oxyalkyl or

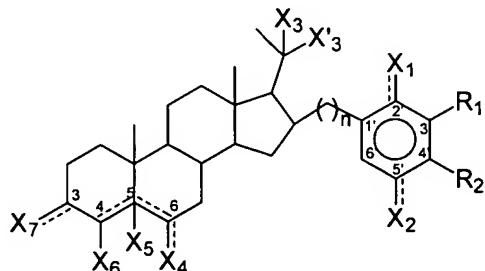
oxycarbonyl, wherein X_3' is glucosyl, fructosyl, galactosyl, mannosyl, fucosyl, cellobiosyl, gentiobiosyl, 2-amino-2-deoxy glucosyl, 2-acetamido-2-deoxy-glucosyl, 2-amino-2-deoxy galactosyl, 2-acetamido-2-deoxy-galactosyl, a disaccharide or a trisaccharide thereof, wherein X_4 is hydrogen, wherein X_5 participates to in a double bond between the carbon atoms in position 5 and 6, wherein X_6 is $-H$, wherein X_7 is hydrogen, oxygen, hydroxyl or oxo, and wherein n is 0.

8. (Currently amended) A-The compound according to ~~any of claims 1 to 4~~claim 1, wherein X_1 and X_2 are $-OMe$, wherein R_1 and R_2 are $-H$, wherein X_3 is glucosyl, fructosyl, galactosyl, mannosyl, fucosyl, cellobiosyl, gentiobiosyl, 2-amino-2-deoxy glucosyl, 2-acetamido-2-deoxy glucosyl, 2-amino-2-deoxy galactosyl, 2-acetamido-2-deoxy-galactosyl, a disaccharide or a trisaccharide thereof, wherein X_3' is hydrogen, alkyl or aralkyl, wherein X_4 is hydrogen, wherein X_5 participates ~~to~~in a double bond between the carbon atoms in position 5 and 6, wherein X_6 is $-H$, wherein X_7 is hydrogen, oxygen, hydroxyl or oxo, and wherein n is 0.

9. (Currently amended) ~~A-The~~ compound according to ~~any of claims 1 to 4~~claim 1, wherein X_1 and X_2 are $-OMe$, wherein R_1 and R_2 are $-H$, wherein X_3 participates together with X_3' ~~to~~in an oxo functional group, wherein X_4 is hydroxyl, glucosyl, fructosyl, galactosyl, mannosyl, fucosyl, cellobiosyl, gentiobiosyl, 2-amino-2-deoxy glucosyl, 2-acetamido-2-deoxy-glucosyl, 2-amino-2-deoxy galactosyl, 2-acetamido-2-deoxy-galactosyl, a disaccharide or a trisaccharide thereof, wherein X_5 participates ~~to~~in a double bond between the carbon atoms in position 5 and 6, wherein X_6 is $-H$, wherein X_7 is hydrogen, oxygen, hydroxyl or oxo, and wherein n is 0.

10. (Currently amended) Compound A compound of formula IB or a pharmaceutically acceptable salt thereof, wherein X_1 , X_2 , X_3 , X_3' , X_4 , X_5 , X_6 , X_7 , R_1 , R_2 and n are selected as indicated in Table A or Table B.

11. Method A method for synthesizing a compound having the structural formula I B

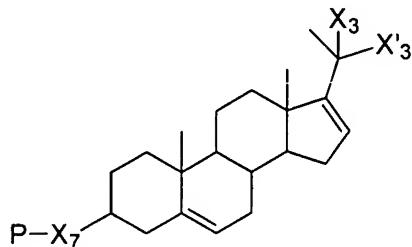


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formula IB

wherein X_1 , X_2 , X_3 , X_4 , X_5 , X_6 , X_7 , R_1 , R_2 and n are selected from the group as indicated in ~~any of claims 1 to 10~~claim 1, said method comprising the steps of

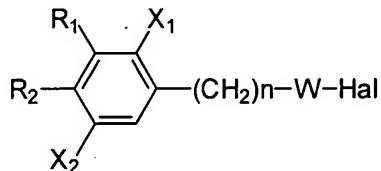
a) providing a starting material having the structural formula IV,



formula IV

wherein X_3 , X_3' and X_7 are selected from the group as indicated in ~~any of claims 1 to 10~~claim 1, and wherein P is a protecting group,

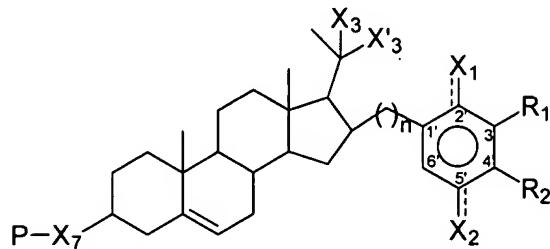
b) effecting reaction between the compound of step a) with an organometallic compound having the structural formula V



formula V

wherein X_1 , X_2 , R_1 , R_2 and n are selected from the group as indicated in ~~any of claims 1 to 10~~claim 1, wherein W is a metal or a combination of metals and wherein Hal is a halogen atom,

to result in an intermediate having the structural formula III'B



formula III'B

wherein X_1 , X_2 , X_3 , X_3' , X_7 , R_1 , R_2 and n are selected from the group as indicated in ~~any of claims 1 to 10~~ claim 1, and wherein p is a protecting group,

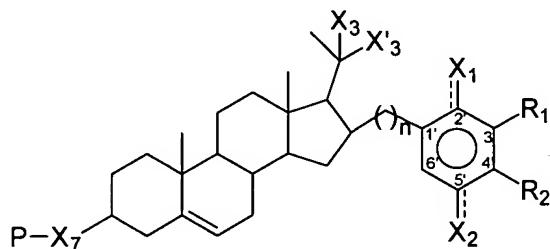
c) effecting reaction between the compound of step b) with an organometallic compound having the structural formula VI



formula VI

wherein $X'3$ is selected from the group as indicated in ~~any of claims 1 to 10~~ claim 1, wherein W is a metal or a combination of metals, and wherein Hal is a halogen atom,

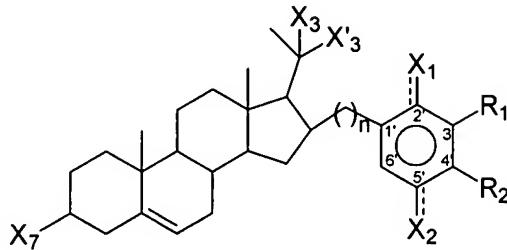
to result in an intermediate having the structural formula IIIB



formula IIIB

wherein X_1 , X_2 , X_3 , X_3' , X_7 , R_1 , R_2 and n are selected from the group as indicated in ~~any of claims 1 to 10~~ claim 1, wherein P is a protecting group,

d) deprotecting the X_7 group of the compound obtained in step c) to form an a compound having the structural formula IIB



formula II B

wherein X_1 , X_2 , X_3 , X_3' , X_7 , R_1 , R_2 and n are selected from the group as indicated in any of claims 1 to 10 claim 1, and

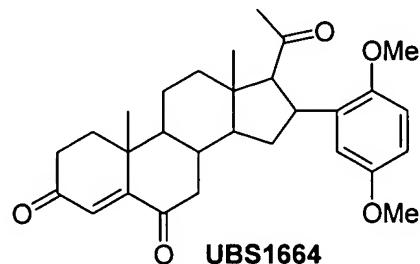
e) oxidizing by reaction with a suitable oxidizing agent or agents to form a compound of formula IB or

e) coupling an O-protected glycosyl or non-protected glycosyl to form a compound of formula IIB wherein X_1 , X_2 , X_3 , X_3' , X_7 , R_1 , R_2 and n are selected from the group as indicated in any of claims 1 to 10 claim 1 and X_7 is an O-protected glycosyl or a non-protected glycosyl, and

f) deprotecting the O-protected groups of glycosyl to form a compound of formula IB wherein X_1 , X_2 , X_3 , X_3' , X_4 , X_5 , X_6 , R_1 , R_2 and n are selected from the group as indicated in any of claims 1 to 10 claim 1, and X_7 is a glycosyl, thio derivatives thereof, amino derivatives thereof, or hydroxyl-protected derivatives thereof.

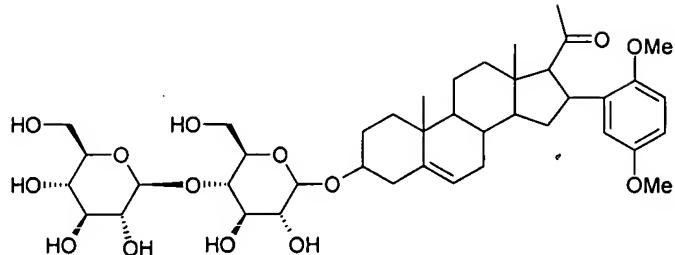
12. (Original) A compound obtainable by any of the steps according to the method of claim 11.

13. (Original) A compound designated as compound UBS1664



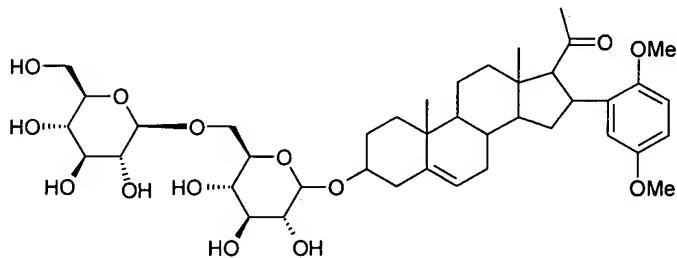
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14. (Original) A compound designated as compound UBS3327.



UBS3327

15. (Original) A compound designated as compound UBS3328.



UBS3328

16. (Cancelled)

17. (Cancelled)

18. (Currently amended) A pharmaceutical composition comprising a pharmaceutically acceptable excipient and a therapeutically effective amount of a compound according to any of claims 1 to 10 and 12 to 15 Claim 1.

19. (Cancelled)

20. (Currently amended) Method A method of treating cancer comprising administrating to an individual in need of such treatment a pharmaceutical composition according to claim 18.

21. (New) A pharmaceutical composition comprising a pharmaceutically acceptable excipient and a therapeutically effective amount of a compound prepared by the method of Claim 12.

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22. (New) A method of treating cancer comprising administrating to an individual in need of such treatment a pharmaceutical composition according to claim 21.